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(54) Title: PRESSURE SENSITIVE ADHESIVE COMPOSITION FOR ELECTRO-MAGNETIC SHIELDING FILM AND
ELECTRO-MAGNETIC SHIELDING FILTER USING THE SAME

(57) Abstract: The present invention relates to an adhesive composition for an electro-magnetic shielding film and an electro-mag-
netic shielding filter using the same, and more particularly to an adhesive composition for an electromagnetic shielding film com-
prising: a) an acrylic copolymer of i) (meth)acrylate ester monomers having C₁ to C₁₂ alkyl groups and ii) vinylic monomers having
hydroxyl groups; and b) a multifunctional isocyanate crosslinking agent. The electro-magnetic shielding filter of the present inven-
tion offers good durability because transparentization is performed with an acrylic adhesive composition having no acidic groups,
and thus oxidation of copper is prevented.

WO 2004/083333 A1

DESCRIPTION

PRESSURE SENSITIVE ADHESIVE COMPOSITION FOR ELECTRO-MAGNETIC SHIELDING FILM AND ELECTRO-MAGNETIC SHIELDING FILTER USING THE SAME

5 **[Technical Field]**

The present invention relates to an adhesive composition for an electro-magnetic shielding film and an electro-magnetic shielding filter using the same, and more particularly to an adhesive composition for an electro-magnetic shielding film wherein the adhesive contains no acidic groups so
10 that oxidation of copper is prevented and thereby having superior durability, and an electro-magnetic shielding filter using the same.

[Background Art]

Recently, with interest in digital televisions and high-quality wide-screen televisions increasing, development of cathode ray tubes (CRTs),
15 liquid crystal displays (LCDs), and plasma display panels (PDPs) is actively progressing.

CRTs, which are widely used as displays in TVs, are superior in resolution and image quality. However, they are not suitable for wide screens of 40 inches or larger because of size and weight.

20 LCDs are superior in terms of power consumption and driving voltage, but they are difficult to make into wide screens and are limited in view angle.

PDPs can be made into wide screens of about 60 inches. When manufacturing a PDP device, a partitioned barrier is formed on a bottom plate and phosphor layers of red, green, and blue are formed on the barrier. The bottom plate is then covered by a top plate, so that electrodes of the bottom plate and those of the top plate face each other in parallel. Then, discharge gas is injected therein and the device is sealed. Light radiated from plasma when the gas is discharged is used to transfer images, and the barrier separates each unit cell.

PDP devices are classified into direct current (DC) and alternating current (AC) types. In DC devices, electrodes are exposed to the discharge area, and in AC devices, a dielectric glass layer is formed on electrodes.

Referring to FIG. 1, a general PDP comprises a case 11, a cover 15 covering said case 11, a drive circuit board 12 located in said case 11, a panel assembly 13, and an electro-magnetic shielding filter 14. Said electro-magnetic shielding filter 14 is coated with a conducting film made of a conducting material, which offers a grounding path passing through the cover 15 to the case 11, and is placed on a glass plate. That is, an electromagnetic wave generated from the panel assembly 13 is passed through the conducting film of the electro-magnetic shielding filter 14 to the cover 15 and the case 11 before it reaches the user. For this purpose, the conducting film of the electro-magnetic shielding filter 14 is electrically

connected to the cover 15 and the case 11.

In addition to shielding electromagnetic waves, said electro-magnetic shielding filter shields near infrared rays, protects the panel from external impact, improves color resolution, prevents reflection, and
5 enhances contrast.

Formerly, transparent conducting films satisfying industrial class A (surface resistance $< 2.5 \Omega/\square$) standard, such as an Ag multi-deposited film, were widely used to manufacture electro-magnetic shielding filter films for plasma display panels. But recently, with the increase in demand for
10 plasma display panels for families, copper etching meshes or conducting fiber meshes that satisfy the class B (surface resistance $< 1.5 \Omega/\square$) standard in which copper or nickel is coated on polyester fiber by electroless plating are being widely used.

Generally, mesh-type electro-magnetic shielding films are
15 manufactured by applying copper foil having a thickness of about $10 \mu\text{m}$ on a polyethylene terephthalate substrate using a transparent adhesive and etching it to pitch $\approx 300 \mu\text{m}$ and linewidth $\approx 10 \mu\text{m}$.

When this electro-magnetic shielding film is laminated with a functional film such as an antireflection film, an aperture is formed between
20 the two films due to the copper foil mesh. Then, because of the difference in refractive indices of air and the polymer layer, light emergent from the plasma display panel becomes scattered, thereby reducing visibility.

Therefore, the aperture should be removed by a process called transparentization.

For transparentization, a method of using a hot melt resin (Japan Patent Publication No. Hei 9-258378), a method of using adhesive, a method of filling a fluid resin (US Patent No. 6,197,408), and so forth have been developed. Among these, the most simple and widely used method is the method of using adhesive by which transparentization can be attained by applying pressure without an additional process. In transparentization using adhesive, the durability of the electro-magnetic shielding filter largely depends on the component of the adhesive.

In general, rubber, acryl, and silicone are widely used for adhesives. Among these, acrylic adhesives are the most popular for preparing functional adhesive compositions because of various melt properties. General acrylic adhesives have superior adhesivity when pressurized slightly at normal temperature. This is because polymer molecules comprising the adhesives are fluid and sensitive to pressure.

The electro-magnetic shielding filter offers colors with good resolution and image quality because of each functional film. If the filter discolours during use, it cannot offer good resolution and image quality. Therefore, there should be no change in transmittivity and color between before and after a durability test. Especially, color change due to oxidation of the copper mesh under high temperature and high humidity is a serious

problem.

The durability of adhesive at the area where it contacts the copper mesh is a problem. Copper looks red in a neutral state, but it turns blue because of Cu^{2+} when oxidized. Therefore, the cause of discoloration of the transparentized adhesive is oxidation of copper.

In widely used acrylic adhesive compositions, several functional groups are used to crosslink the adhesive. Particularly, when monomers having acidic groups ($-\text{COOH}$) are used, the acidic groups ($-\text{COOH}$) oxidize copper under high temperature and high humidity, thereby discoloring the electro-magnetic shielding filter and reducing color resolution.

[Description of the drawings]

FIG. 1 shows the structure of a plasma display panel.

FIGs. 2a and 2b show cross-sections of an electro-magnetic shielding film before and after transparentization.

FIG. 3 shows the cross-section of a electro-magnetic shielding filter of a plasma display panel.

FIGs. 4a to 4d show transmittivity measurement results for Examples 1 to 3 and Comparative Example 1 before and after a durability test.

Description of marks for a main part of drawings

11 --- case

12 --- drive circuit board

13 --- panel assembly

14 --- electro-magnetic shielding filter

- 15 — cover
21, 31 — transparent substrate
22, 32 — adhesive layer
23, 33 — layers of copper foil
24, 34 — black screen
25, 35 — adhesive
26, 36 — transparent substrate
27, 37 — reinforcing glass
5 38 — Color correction layer
39 — near infrared ray shielding layer
40 — near infrared ray shielding layer

[Technical Problem]

It is an object of the present invention to provide an adhesive composition for an electro-magnetic shielding film having superior durability by preventing oxidization of copper, and an electro-magnetic shielding filter using the same.

[Technical Solution]

To attain said object, the present invention provides an acrylic adhesive composition for an electro-magnetic shielding film, comprising:

- 15 a) an acrylic copolymer of i) (meth)acrylate ester monomers having C₁ to C₁₂ alkyl groups; and ii) vinylic monomers having hydroxyl groups; and
b) a multifunctional isocyanate crosslinking agent.

The present invention also provides an electro-magnetic shielding filter for a plasma display panel using said acrylic adhesive composition for
20 an electro-magnetic shielding film.

[Advantageous Effects]

Because the electro-magnetic shielding filter of the present invention is transparentized with an acrylic adhesive composition having no acidic groups, oxidation of copper is prevented, which improves durability and prevents discoloration of the electro-magnetic shielding filter.

[Best Mode]

Hereinafter, the present invention is described in more detail referring to the attached drawings.

As seen in FIG. 2a, an electro-magnetic shielding film before transparentization comprises layers of copper foil 23, black screen 24, adhesive 25, and transparent substrate 26 from the top side down. An air layer is formed when the transparent substrate 21 such as an antireflection film is glued to the copper foil 23. The difference in refractive indices of this air layer and the transparent substrate 21, which is made of polymer film, causes scattering of light emergent from the plasma display panel, thereby reducing visibility.

Accordingly, a crosslinking agent, UV or EB, is used to crosslink the adhesive composition and therefore remove the air layer, as seen in FIG. 2b. This process of removing the air layer is called transparentization.

If the adhesive used in the adhesive composition for the electro-magnetic shielding film of the present invention has too low elasticity because of too low crosslinking density or molecular weight, air bubbles

may form at high temperature, which may cause scattering. Otherwise, if an adhesive having too large elasticity is used for a long time, excessive crosslinkage may cause peeling of layers. Since viscoelasticity of an adhesive is mainly dependent on molecular weight, molecular weight
5 distribution, or molecular structure of the polymer chain, and particularly molecular weight, it is preferable to use an acrylic polymer having a molecular weight ranging from 300,000 to 2,000,000. Said adhesive may be prepared by conventional radical copolymerization.

To be more specific, the acrylic adhesive composition for the electro-
10 magnetic shielding film preferably comprises:

a) 100 parts by weight of an acrylic copolymer of:

i) 90 to 99.9 parts by weight of (meth)acrylate ester monomers having C_1 to C_{12} alkyl groups; and

ii) 0.1 to 10 parts by weight of vinylic monomers having
15 hydroxyl groups; and

b) 0.01 to 10 parts by weight of a multifunctional isocyanate crosslinking agent.

More preferably, said (meth)acrylate ester monomers having C_1 to C_{12} alkyl groups are comprised at 94 to 99.9 parts by weight for 100 parts
20 by weight of said acrylic copolymer. If the content is below 90 parts by weight, initial adhesivity becomes poor and cost increases.

Preferably, for said (meth)acrylate ester monomers having C_1 to C_{12}

alkyl groups; an alkyl ester of acrylate (or methacrylate) having one to twelve carbon atoms and preferably having two to eight carbon atoms, is used. Preferably, said alkyl(meth)acrylate has two to eight carbon atoms in the alkyl groups because cohesiveness of the adhesive reduces if the alkyl groups are too long. To be specific, butyl(meth)acrylate, 2-ethylhexyl(meth)acrylate, ethyl(meth)acrylate, methyl(meth)acrylate, *n*-propyl(meth)acrylate, isopropyl(meth)acrylate, *t*-butyl(meth)acrylate, pentyl(meth)acrylate, *n*-octyl(meth)acrylate, or isononyl(meth)acrylate may be used.

Also, such monomers as acrylonitrile, styrene, glycidyl(meth)acrylate, and vinyl acetate may be used to control the glass transition temperature of the adhesive or offer other properties.

Said vinylic monomers having hydroxyl groups are used to offer cohesiveness, so that the adhesive is not damaged at high temperature, through reaction with said multifunctional isocyanate crosslinking agent. Preferably, said vinylic monomers are comprised at 0.1 to 10 parts by weight for 100 parts by weight of said acrylic copolymer. If the content is below 0.1 part by weight, the adhesive may be easily damaged at high temperature. Otherwise, if it exceeds 10 parts by weight, fluidity at high temperature may decrease.

For said vinylic monomers having hydroxyl groups, 2-hydroxyethyl(meth)acrylate, 2-hydroxypropyl(meth)acrylate, 2-

hydroxyethyleneglycol(meth)acrylate, or 2-
hydroxypropyleneglycol(meth)acrylate may be used.

Preferably, said multifunctional isocyanate crosslinking agent is
comprised at 0.01 to 10 parts by weight for 100 parts by weight of said
5 acrylic copolymer. If the content is below 0.01 part by weight, crosslinkage
becomes insufficient. Otherwise, if it exceeds 10 parts by weight, peeling
of layers may occur due to excessive crosslinkage.

Said multifunctional isocyanate crosslinking agent maintains
cohesiveness of the adhesive at high temperature through crosslinkage,
10 and therefore improves adhesion reliability. For said multifunctional
isocyanate crosslinking agent, tolylene diisocyanate, diphenylmethane
diisocyanate, hexamethylene diisocyanate, or a trimethylolpropane adduct
of tolylene diisocyanate may be used. Said multifunctional isocyanate
crosslinking agent does not cause crosslinkage of isocyanate groups during
15 formation of the adhesive layer, which enables uniform coating. After the
coating process is completed, a crosslinkage is formed while passing
through drying and aging processes. Resultantly, an adhesive layer with
improved cohesiveness is obtained. The improved cohesiveness improves
physical properties and cuttability of the adhesive.

20 Also, a silane coupling agent may be added to improve adhesion
durability. Said silane coupling agent improves adhesivity with time and
heat, and thereby prevents air bubbling and peeling under high temperature

and high humidity, and enhances durability and reliability. Preferably, said silane coupling agent is comprised at 0.01 to 10 parts by weight for 100 parts by weight of said acrylic copolymer. If the content is below 0.01 part by weight, improvement of adhesivity with time and heat is only slight.

5 Otherwise, if it exceeds 10 parts by weight, air bubbling or peeling may occur, which reduces durability and reliability.

For said silane coupling agent, β -(3,4-epoxycyclohexyl) ethyltrimethoxysilane, γ -glycidoxypropyltrimethoxysilane, γ -glycidoxypropylmethyldiethoxysilane or γ -glycidoxypropyltriethoxysilane
10 may be used.

In addition, 1 to 100 parts by weight of an adhesive resin may be used to improve adhesivity. However, if said adhesive resin is used excessively, cohesiveness of the adhesive may be reduced. For said adhesive resin, (hydrated) hydrocarbon resin, (hydrated) rosin resin, (hydrated) rosin ester resin, (hydrated) terpene resin, (hydrated) terpene
15 phenol resin, polymeric rosin resin, or polymeric rosin ester resin may be used.

In the present invention, said acrylic copolymer may be polymerized by solution polymerization, photopolymerization, bulk polymerization, suspension polymerization, or emulsion polymerization, and preferably by
20 solution polymerization. Preferably, the polymerization temperature is 50 to 140°C, and an initiator is added when the monomers are mixed uniformly.

Preferably, the acrylic adhesive resin for the electro-magnetic shielding film of the present invention has a crosslinking density ranging from 1 to 95%.

The acrylic adhesive resin for the electro-magnetic shielding film composition of the present invention may be prepared by well-known photopolymerization curing by selecting an appropriate photoinitiator. The acrylic adhesive resin composition of the present invention can be used for polarizers, optical functional films, optical control films, etc. Also, it can be used for industrial sheets including reflective sheets, structural adhesive sheets, photographic adhesive sheets, adhesive sheets for drawing traffic lanes, optical adhesives, and adhesives for electronic devices.

During transparentization of the acrylic adhesive composition for the electro-magnetic shielding film of the present invention, it is preferable to dilute the acrylic adhesive composition with an organic solvent before coating to improve its coating characteristics.

Preferably, a multifunctional crosslinking agent is added as the crosslinking agent. More preferably, an isocyanate crosslinking agent, epoxy crosslinking agent, aziridine crosslinking agent, or metal chelate crosslinking agent is added. For said isocyanate crosslinking agent, tolylene diisocyanate, diphenylmethaneisocyanate, or hexamethylene diisocyanate may be used.

Said multifunctional isocyanate crosslinking agent does not cause

crosslinkage of isocyanate groups during formation of the adhesive layer, which enables uniform coating. After the coating process is completed, a crosslinkage is formed while passing through drying and aging processes. Resultantly, an adhesive layer with improved cohesiveness is obtained.

- 5 The improved cohesiveness improves such physical properties as durability, reliability, adhesivity, and cuttability of the adhesive.

The crosslinking may be attained using UV or EB, not using a crosslinking agent.

- 10 The acrylic adhesive composition for the electro-magnetic shielding film of the present invention may further comprise a near infrared ray absorbent, epoxy resin, curing agent, silane coupling agent, plasticizer, UV stabilizer, antioxidant, dye, reinforcing agent, or filler, if required.

- 15 Durability of the electro-magnetic shielding film after transparentization is tested by exposing it under a high temperature of 80°C for 500 hours or under 60°C and RH (relative humidity) of 90% for 500 hours, and measuring transmittivity change before and after exposure. The larger the transmittivity change, the less the durability.

- 20 If transparentization is performed using the adhesive prepared by using an acrylic copolymer having acidic groups, the residual acidic groups oxidize copper, which turns blue. Resultantly, transmittivity of the electro-magnetic shielding film is significantly reduced and durability of the electro-magnetic shielding film is reduced as well.

However, in the present invention, the air layer is removed by crosslinkage using the adhesive composition for the electro-magnetic shielding film. Therefore, the transparentization process is simple. Also, because acryl monomers having no acidic groups are used, oxidation of copper is prevented after crosslinkage. Accordingly, transmittivity of the electro-magnetic shielding film changes little even under high temperature or high humidity, thereby improving durability.

The present invention also provides a plasma display panel electro-magnetic shielding filter manufactured by laminating layers of reinforcing glass 37, a color correction layer 38, and a near infrared ray shielding layer 39 below the transparent substrate 36 of the electro-magnetic shielding film, on which layers of copper foil 33, black screen 34, and adhesive 35 are laminated, and laminating antireflection films 40 below the near infrared ray shielding layer 39 and on top of the adhesive layer 32. Another transparent substrate 31 may be laminated below the antireflection film 40.

Because vinylic monomers having hydroxyl groups but having no acidic groups are used in the adhesive composition for the electro-magnetic shielding filter of the present invention, oxidation of copper is prevented even under high temperature and high humidity, which reduces transmittivity of the electro-magnetic shielding filter and improves durability of the plasma display panel.

Hereinafter, the present invention is described in more detail through

examples and a comparative example. However, the following examples and comparative example are only for the understanding of the present invention, and the present invention is not limited by the following examples.

Example 1

5 (Preparation of acrylic copolymer)

A monomer mixture comprising 98 parts by weight of *n*-butylacrylate (BA) and 2 parts by weight 2-hydroxyethyl(meth)acrylate (2-HEMA) was put in a 1,000 cc reactor equipped with nitrogen reflux and cooling units. Then, 100 parts by weight of ethyl acetate (EAc) were added as a solvent. After
10 the reactor was purged with nitrogen gas for 20 minutes to remove oxygen, the temperature was set at 60°C. Then, 0.03 parts by weight of azobisisobutyronitrile (AIBN) diluted to 50% in ethyl acetate was added as an reaction initiator. Reaction was performed for 8 hours to prepare an acrylic copolymer (PA-1) having a molecular weight of 1,500 K (measured
15 with standard polystyrene sample).

(Preparation of acrylic adhesive composition)

1.5 parts by weight of a tolylene diisocyanate adduct of trimethylolpropane (TDI-1) diluted to 10 wt% in ethyl acetate was added for 100 parts by weight of the prepared acrylic copolymer (PA-1) as a
20 multifunctional isocyanate crosslinking agent. The resultant mixture was diluted to an appropriate concentration considering the coating characteristics, and mixed uniformly. Then, it was coated on a 38 µm-thick

releasing film and dried to prepare a uniform adhesive layer measuring 30 μm .

Transparentization of the electro-magnetic shielding film was performed using the prepared adhesive layer.

5 Example 2

The procedure of Example 1 was performed except for using 2-hydroxypropyl(meth)acrylate instead of 2-hydroxyethyl(meth)acrylate (2-HEMA).

Example 3

10 (Preparation of acrylic copolymer)

An acrylic copolymer (PA-2) having a molecular weight of 1,000 K (measured with standard polystyrene sample) was prepared in the same manner of Example 1 using a monomer mixture comprising 94 parts by weight of *n*-butylacrylate (BA) and 6 parts by weight of 2-hydroxyethyl(meth)acrylate (2-HEMA), and 0.04 part by weight of azobisisobutyronitrile (AIBN) diluted to 50% in ethyl acetate as a reaction initiator.

(Preparation of acrylic adhesive composition)

The procedure of Example 1 was performed except for using PA-2 as acrylic copolymer.

Comparative Example 1

(Preparation of acrylic copolymer)

A monomer mixture comprising 94 parts by weight of *n*-butylacrylate (BA) and 6 parts by weight acrylate (acetic acid) was put in a 1,000 cc reactor equipped with nitrogen reflux and cooling units. Then, 100 parts by weight of ethyl acetate (EAc) were added as solvent. After the reactor was purged with nitrogen gas for 20 minutes to remove oxygen, the temperature was set at 60°C. Then, 0.04 parts by weight of azobisisobutyronitrile (AIBN) diluted to 50% in ethyl acetate was added as a reaction initiator.

10 The reaction was performed for 8 hours to prepare an acrylic copolymer (PA-3) having a molecular weight of 1,000 K (measured with standard polystyrene sample).

(Preparation of acrylic adhesive composition)

1.5 parts by weight of a tolylene diisocyanate adduct of trimethylolpropane (TDI-1) diluted to 10 wt% in ethyl acetate was added for 100 parts by weight of the prepared acrylic copolymer (PA-3) as a multifunctional isocyanate crosslinking agent. The resultant mixture was diluted to an appropriate concentration considering the coating characteristics, and mixed uniformly. Then, it was coated on a 38 μm -thick releasing film and dried to prepare a uniform adhesive layer measuring 30 μm .

Transparentization of the electro-magnetic shielding film was

performed using the prepared adhesive layer.

(Evaluation of durability)

Each adhesive composition prepared in Examples 1 to 3 and Comparative Example 1 was treated on an electro-magnetic shielding film.

5 After transparentization, transmittivity was measured in a wavelength range of 300 nm to 900 nm. After exposure under 60°C and RH 90% for 500 hours, transmittivity was measured again. The transmittivity measurement results are shown in FIGs. 4a to 4d. Transmittivity change at 680 nm is shown in the following Table 1.

10 Table 1

	Transmittivity before exposure	Transmittivity after exposure	Transmittivity change (%)
Example 1	76.5	76.5	0
Example 2	80.6	80.1	0.5
Example 3	79.4	79.0	0.4
Comparative Example 1	76.5	67.3	9.2

As seen in FIGs. 4a to 4d and Table 1, electro-magnetic shielding films transparentized with acrylic adhesive compositions of Example 1 to 3, which have no acidic groups, showed little transmittivity change before and
15 after exposure. On the other hand, the electro-magnetic shielding film

transparentized with the adhesive composition of Comparative Example 1, which has acidic groups (-COOH), experienced a transmittivity change of 9.2%.

From this result, it can be seen that oxidation of copper in the copper
5 foil is prevented if transparentization is performed using the acrylic adhesive composition of the present invention, which has no acidic groups, and therefore there is little transmittivity change even under high temperature and high humidity, thereby improving durability.

While the present invention has been described in detail with
10 reference to the preferred embodiments, those skilled in the art will appreciate that various substitutions and modifications can be made thereto without departing from the spirit and scope of the present invention as set forth in the appended claims.

What is claimed is:

1. An acrylic adhesive composition for an electro-magnetic shielding film, comprising:

- a) an acrylic copolymer of i) (meth)acrylate ester monomers having C₁ to C₁₂ alkyl groups and ii) vinylic monomers having hydroxyl groups; and
b) a multifunctional isocyanate crosslinking agent.

2. The acrylic adhesive composition for an electro-magnetic shielding film of Claim 1, which comprises:

- a) 100 parts by weight of an acrylic copolymer of:
i) 90 to 99.9 parts by weight of (meth)acrylate ester monomers having C₁ to C₁₂ alkyl groups; and
ii) 0.1 to 10 parts by weight of vinylic monomers having hydroxyl groups; and
b) 0.01 to 10 parts by weight of a multifunctional isocyanate crosslinking agent.

3. The acrylic adhesive composition for an electro-magnetic shielding film of Claim 1, wherein said (meth)acrylate ester monomers having C₁ to C₁₂ alkyl groups are one or more members selected from the group consisting of butyl(meth)acrylate, 2-ethylhexyl(meth)acrylate, ethyl(meth)acrylate, methyl(meth)acrylate, *n*-propyl(meth)acrylate, isopropyl(meth)acrylate, *t*-butyl(meth)acrylate, pentyl(meth)acrylate, *n*-octyl(meth)acrylate, and isononyl(meth)acrylate.

4. The acrylic adhesive composition for an electro-magnetic shielding film of Claim 1, wherein said vinylic monomers having hydroxyl groups are one or more members selected from the group consisting of 2-hydroxyethyl(meth)acrylate, 2-hydroxypropyl(meth)acrylate, 2-hydroxyethyleneglycol(meth)acrylate, and 2-hydroxypropyleneglycol(meth)acrylate.

5. The acrylic adhesive composition for an electro-magnetic shielding film of Claim 1, wherein said vinylic monomers having hydroxyl groups react with isocyanate groups of said multifunctional isocyanate crosslinking agent to form a partial crosslinkage.

6. The acrylic adhesive composition for an electro-magnetic shielding film of Claim 1, wherein said multifunctional isocyanate crosslinking agent is one or more members selected from the group consisting of tolylene diisocyanate, diphenylmethane diisocyanate, hexamethylene diisocyanate, and a trimethylolpropane adduct of tolylene diisocyanate.

7. The acrylic adhesive composition for an electro-magnetic shielding film of Claim 1, which further comprises one or more additives selected from the group consisting of a near infrared ray absorbent, epoxy resin, curing agent, silane coupling agent, plasticizer, UV stabilizer, antioxidant, dye, reinforcing agent, and filler.

8. The acrylic adhesive composition for an electro-magnetic

shielding film of Claim 1, wherein said acrylic copolymer is prepared by solution polymerization, photopolymerization, bulk polymerization, suspension polymerization, or emulsion polymerization.

9. The acrylic adhesive composition for an electro-magnetic
5 shielding film of Claim 1, which has a crosslinking density ranging from 1 to 95%.

10. An electro-magnetic shielding filter for a plasma display panel transparentized using an acrylic adhesive composition of any of Claims 1 to 9.

[received by the International Bureau on 30 August 2004 (30.08.2004);
original claims 1 and 10 amended; original claim 9 cancelled;
remaining claims unchanged (3 pages)]

1. (Amended) An acrylic adhesive composition for an electro-magnetic
shielding film having crosslinking density ranging from 1 to 95%, comprising:

- a) an acrylic copolymer of i) (meth)acrylate ester monomers having
5 C₁ to C₁₂ alkyl groups and ii) vinyllic monomers having hydroxyl groups; and
b) a multifunctional isocyanate crosslinking agent.

2. The acrylic adhesive composition for an electro-magnetic
shielding film of Claim 1, which comprises:

- a) 100 parts by weight of an acrylic copolymer of:
10 i) 90 to 99.9 parts by weight of (meth)acrylate ester
monomers having C₁ to C₁₂ alkyl groups; and
ii) 0.1 to 10 parts by weight of vinyllic monomers having
hydroxyl groups; and
b) 0.01 to 10 parts by weight of a multifunctional isocyanate
15 crosslinking agent.

3. The acrylic adhesive composition for an electro-magnetic
shielding film of Claim 1, wherein said (meth)acrylate ester monomers
having C₁ to C₁₂ alkyl groups are one or more members selected from the
group consisting of butyl(meth)acrylate, 2-ethylhexyl(meth)acrylate,
20 ethyl(meth)acrylate, methyl(meth)acrylate, *n*-propyl(meth)acrylate,
isopropyl(meth)acrylate, *t*-butyl(meth)acrylate, pentyl(meth)acrylate, *n*-
octyl(meth)acrylate, and isononyl(meth)acrylate.

4. The acrylic adhesive composition for an electro-magnetic shielding film of Claim 1, wherein said vinylic monomers having hydroxyl groups are one or more members selected from the group consisting of 2-hydroxyethyl(meth)acrylate, 2-hydroxypropyl(meth)acrylate, 2-hydroxyethyleneglycol(meth)acrylate, and 2-hydroxypropyleneglycol(meth)acrylate.

5. The acrylic adhesive composition for an electro-magnetic shielding film of Claim 1, wherein said vinylic monomers having hydroxyl groups react with isocyanate groups of said multifunctional isocyanate crosslinking agent to form a partial crosslinkage.

6. The acrylic adhesive composition for an electro-magnetic shielding film of Claim 1, wherein said multifunctional isocyanate crosslinking agent is one or more members selected from the group consisting of tolylene diisocyanate, diphenylmethane diisocyanate, hexamethylene diisocyanate, and a trimethylolpropane adduct of tolylene diisocyanate.

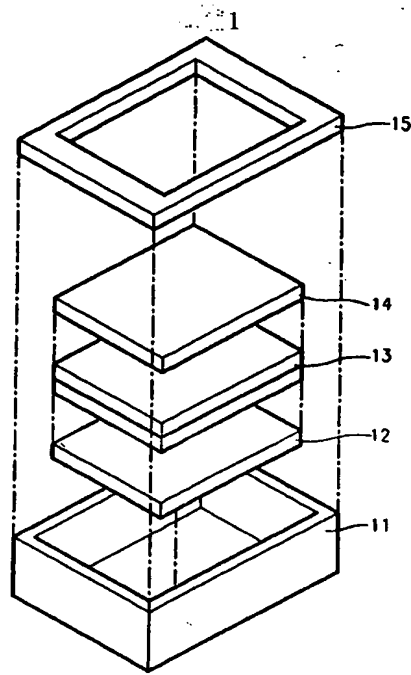
7. The acrylic adhesive composition for an electro-magnetic shielding film of Claim 1, which further comprises one or more additives selected from the group consisting of a near infrared ray absorbent, epoxy resin, curing agent, silane coupling agent, plasticizer, UV stabilizer, antioxidant, dye, reinforcing agent, and filler.

8. The acrylic adhesive composition for an electro-magnetic

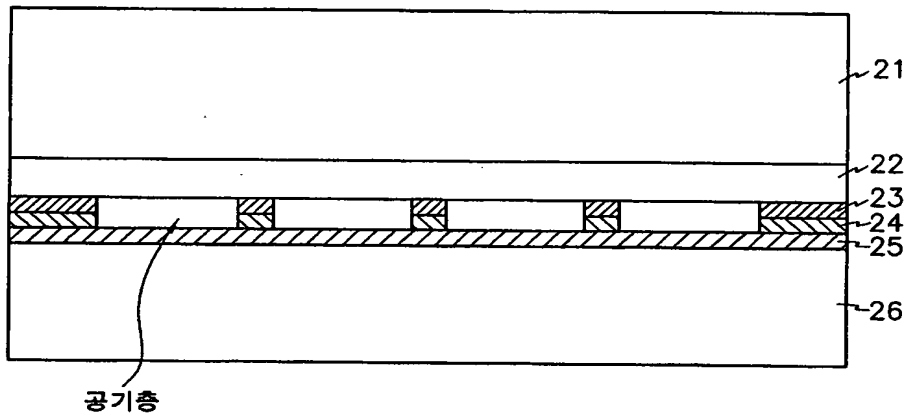
shielding film of Claim 1, wherein said acrylic copolymer is prepared by solution polymerization, photopolymerization, bulk polymerization, suspension polymerization, or emulsion polymerization.

9. (Cancelled)

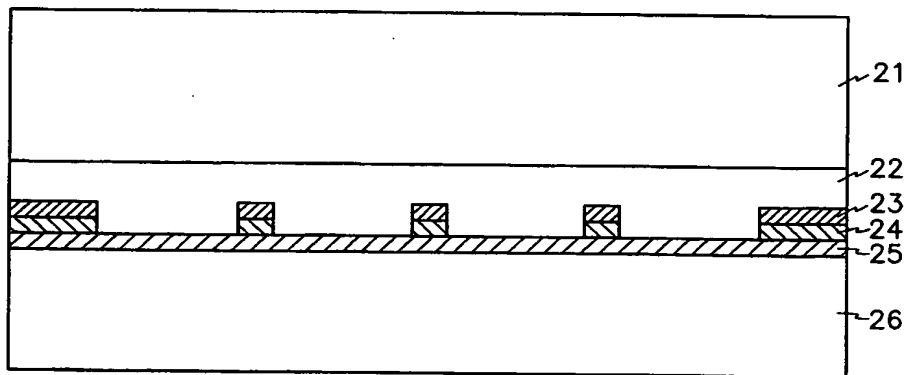
- 5 10. (Amended) An electro-magnetic shielding filter for a plasma display panel transparentized using an acrylic adhesive composition of any of Claims 1 to 8.



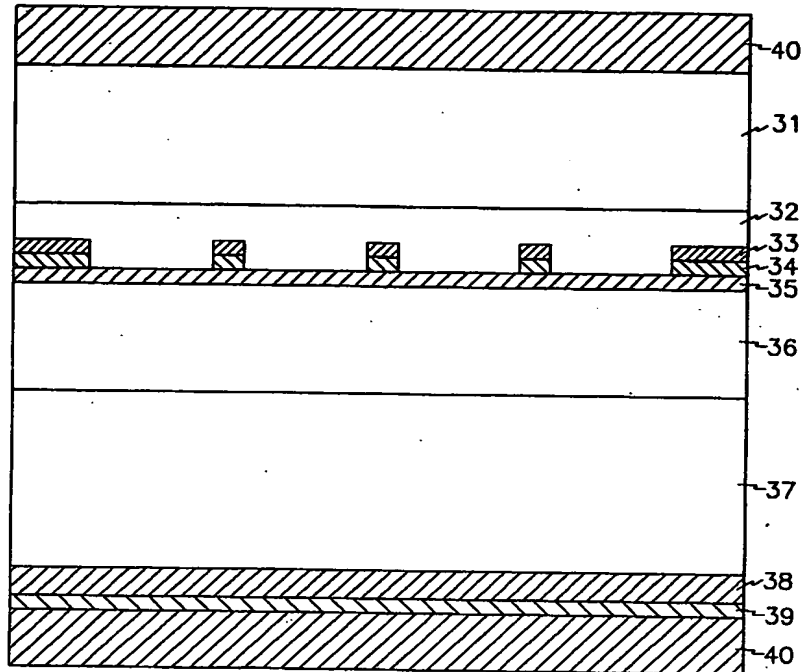
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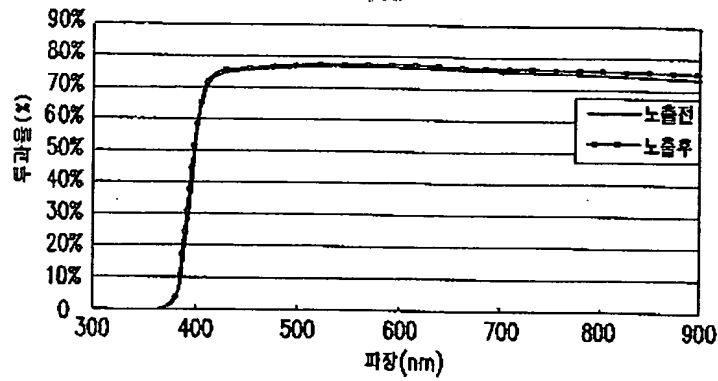
2b



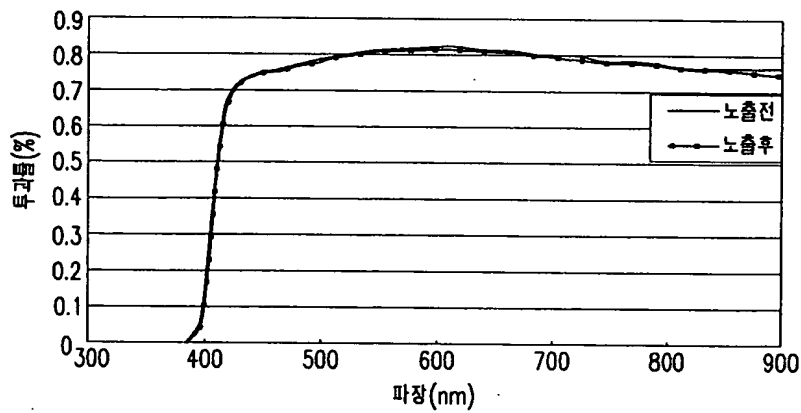
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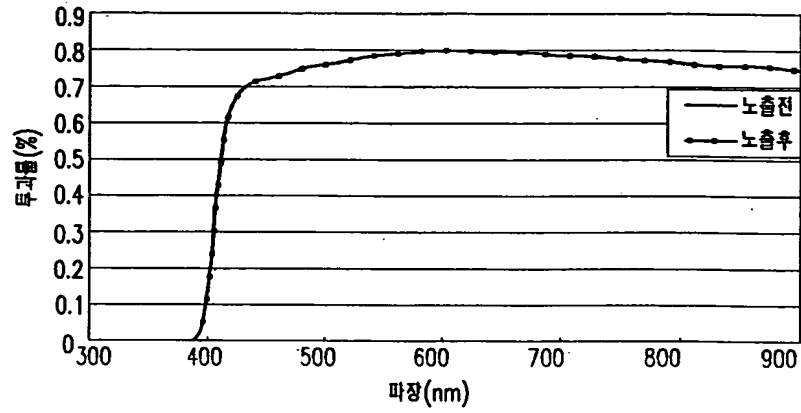
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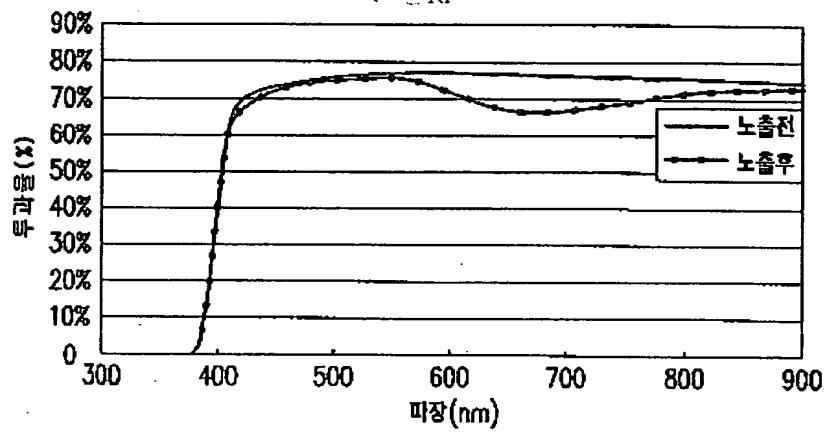
4b



14c





14d



INTERNATIONAL SEARCH REPORT

International application No.

KR2004/000544

A. CLASSIFICATION OF SUBJECT MATTER IPC7 C09J 133/10 According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) IPC7 C09J 7/02, C09J 133/04, C09J 175/04 Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) PAJ, E-SPACENET, E-KIPASS		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	JP 09-040927 A (NIPPON CARBIDE IND CO INC) 10 FEBRUARY 1997 See claims 1-5	1-10
Y	JP 59-152977 A (NITTO ELECTRIC IND CO LTD) 31 AUGUST 1984 See patent abstract	1-10
A	JP 10-152658 A (PORATEKUNO KK) 9 JUNE 1998 See the whole document	1
A	JP 08-081658 (TERAOKA SEISAKUSHO KK ET AL) 26 MARCH 1996 See the whole document	1
P, Y	JP 2003-119446 A (NIPPON CARBIDE IND CO INC) 23 APRIL 2003 See claims 1-10	1-10
<input type="checkbox"/> Further documents are listed in the continuation of Box C. <input checked="" type="checkbox"/> See patent family annex.		
<p>* Special categories of cited documents:</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier application or patent but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</p> <p>"&" document member of the same patent family</p>		
Date of the actual completion of the international search 30 JUNE 2004 (30.06.2004)		Date of mailing of the international search report 30 JUNE 2004 (30.06.2004)
Name and mailing address of the ISA/KR  Korean Intellectual Property Office 920 Dunsan-dong, Seo-gu, Daejeon 302-701, Republic of Korea Facsimile No. 82-42-472-7140		Authorized officer KIM, Seung Soo Telephone No. 82-42-481-5581 

INTERNATIONAL SEARCH REPORT
Information on patent family members

International application No.

PCT/KR2004/000544

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JP 59-152977 A	31.08.84	NONE	
JP 10-152658 A	09.06.98	NONE	
JP 08-081658 A	26.03.96	NONE	
JP 2003-119446 A	23.04.03	NONE	